AMENDMENTS TO THE CLAIMS

The following listing of claims replaces all prior versions of claims in the application.

1. (Currently Amended) A method of separating a negatively charged target biopolymer from other biopolymers which are not negatively charged or which are larger than said target biopolymer, comprising the steps of:

partitioning a <u>container into a first solution</u> <u>buffer chamber</u>, <u>initially</u> containing said target biopolymer and other biopolymers, and a second <u>solution</u> <u>buffer chamber</u>, for preserving separated target biopolymer, with the use of a partition;

moving said target biopolymer from within said first solution buffer chamber through said partition into said second solution buffer chamber using electrophoresis; and

separating said target biopolymer from said a buffer in said second solution buffer chamber,

wherein said partition is a gel, a pillar array or a porous filter, wherein said target biopolymer is a nucleic acid or protein, and wherein said other biopolymers are nucleic acids and/or proteins.

2. (Currently Amended) A method of separating a negatively charged target biopolymer from other biopolymers which are smaller than said target biopolymer, comprising the steps of:

partitioning a container into a first solution <u>buffer chamber</u>, <u>initially</u> containing said target biopolymer and said other biopolymers, a second solution <u>buffer chamber</u>, for preserving said other biopolymers, and a third <u>solution</u> <u>buffer chamber</u>, for preserving said target biopolymer,

from each other with the use of a partition;

moving said other biopolymers from within said first solution <u>buffer chamber</u> through said partition and into said second solution <u>buffer chamber using a first electrophoresis</u> device,

moving said target biopolymer from within said first solution buffer chamber into said partition using said first electrophoresis device, then

moving said target biopolymer from within said partition into said third solution buffer chamber using a second electrophoresis device; and

separating said target biopolymer from a buffer in said third solution buffer chamber, wherein said target biopolymer is a nucleic acid or protein, and wherein said other biopolymers are nucleic acids and/or proteins.

- 3. (Previously Presented) The biopolymer separation method of claim 2, wherein said partition is a gel, a pillar array or a porous filter.
- 4. (Withdrawn/Currently Amended) A biopolymer separation apparatus, wherein a negatively charged target biopolymer is separated from among biological samples, comprising:
 - a first solution containing said biological samples;
 - a second solution for preserving separated biopolymers;
- an electrophoresis container carrying a partition to partition said first solution from said second solution;

first positive and negative electrodes provided to move said negatively charged

biopolymer from within said first solution through said partition into said second solution using

electrophoresis; and

a power supply for applying positive and negative voltages to said positive and negative

electrodes respectively,

wherein biopolymer separation can be performed by applying voltages to said electrodes

and moving said target biopolymer from within said first solution through said partition to said

second solution.

5. (Withdrawn/Currently Amended) The biopolymer separation apparatus of claim 4, wherein a

third solution is carried in said container in order to contact said partition in a direction different

from directions of said first solution and said second solution and to preserve said biopolymer

moved through said partition, comprising:

second positive and negative electrodes for electrophoresis which are provided to move

said negatively charged biopolymer from said partition into said third solution using

electrophoresis; and

a power supply for applying positive and negative voltages to said positive and negative

electrodes respectively,

wherein biopolymer separation can be performed by moving said target biopolymer into

said second or third chamber through the switching of movement directions caused by

electrophoresis.

Page 4

6. (Withdrawn) The biopolymer separation apparatus of claim 4 or claim 5, wherein said partition

is a gel, a pillar array or a porous filter.

7. (Currently Amended) A biopolymer separation method, wherein a negatively charged target

biopolymer fixed to a magnetic bead is separated from other biopolymers, comprising the steps

of:

partitioning a container into a first solution buffer chamber, initially containing said target

biopolymer fixed to said magnetic bead and said other biopolymers, a second solution buffer

chamber, for preserving separated other biopolymers, and a third solution buffer chamber, for

preserving said separated target biopolymer fixed to said magnetic bead, from each other with the

use of a partition;

moving said target biopolymer fixed to said magnetic bead and said other biopolymers

from within said first solution buffer chamber into said partition using electrophoresis;

while said target biopolymer fixed to said magnetic bead and said other biopolymers are in

said partition, moving said target biopolymer fixed to said magnetic bead [[,]] into said third

solution buffer chamber using magnetophoresis; and

separating said target biopolymer fixed to said magnetic bead from a buffer in said third

solution buffer chamber,

wherein said target biopolymer is a nucleic acid or protein, and

wherein said other biopolymers are nucleic acids and/or proteins.

Page 5

Amendment under 37 CFR 1.114 Serial No. 10/801,544 Attorney Docket No. 042187

- 8. (Previously Presented) The biopolymer separation method of claim 7, wherein said partition is a gel, a pillar array or a porous filter.
- 9. (Withdrawn) A biopolymer separation apparatus, wherein a negatively charged target biopolymer fixed to a magnetic bead is separated from among biological samples, comprising:
 - a first solution containing said biological samples;
 - a second solution for preserving separated biopolymers;
 - a third solution for preserving a separated target biopolymer fixed to a magnetic bead;
 - a container carrying a partition to partition these three solutions from each other;

positive and negative electrodes provided in said container to move negatively charged biopolymers from within said first solution into said partition and said second solution using electrophoresis;

a power supply to apply positive and negative voltages to said positive and negative electrodes respectively; and

a magnetic field generation means wherein a magnetic field is generated in order to move said target biopolymer fixed to a magnetic bead, which is in transit in said partition using electrophoresis, into said third solution using magnetophoresis,

wherein biopolymer separation can be performed by moving said target biopolymer fixed to a magnetic bead into said third solution using electrophoresis and magnetophoresis.

10. (Withdrawn) The biopolymer separation apparatus of claim 9, wherein said partition is a

Amendment under 37 CFR 1.114 Serial No. 10/801,544 Attorney Docket No. 042187

gel, a pillar array or a porous filter.

11. (Withdrawn) The biopolymer separation apparatus of claim 9 or claim 10, wherein an electromagnet, an electromagnetic coil, or a permanent magnet is used as said magnetic field generation means.